California Medical Device Recall Information

Recall Name
Ventlab Corporation Recalls Manual Resuscitators Due to Possible Malfunction

Recall Date	Product Description	Recalling Firm	Recall Reason
7/11/12	Manual Resuscitators	**Ventlab, Corp.** Mocksville, NC	*Potentially delivers little to no air/oxygen through the patient valve to the patient*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Ventlab Manual Resuscitators Suspect Lots Recalled: • Product List	**CA**, nationwide	Distributed between March 2012 to July 2012

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm324561.htm?source=govdelivery